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NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2	Apr 08	"Ask CAS" for self-help around the clock
NEWS	3	Apr 09	BEILSTEIN: Reload and Implementation of a New Subject Area
NEWS	4	Apr 09	ZDB will be removed from STN
NEWS	5	Apr 19	US Patent Applications available in IFICDB, IFIPAT, and
IFIUDB			
NEWS	6	Apr 22	Records from IP.com available in CAPLUS, HCAPLUS, and
ZCAPLUS			
NEWS	7	Apr 22	BIOSIS Gene Names now available in TOXCENTER
NEWS	8	Apr 22	Federal Research in Progress (FEDRIP) now available
NEWS	9	Jun 03	New e-mail delivery for search results now available
NEWS	10	Jun 10	MEDLINE Reload
NEWS	11	Jun 10	PCTFULL has been reloaded
NEWS	12	Jul 02	FOREGE no longer contains STANDARDS file segment
NEWS	13	Jul 22	USAN to be reloaded July 28, 2002; saved answer sets no longer valid
NEWS	14	Jul 29	Enhanced polymer searching in REGISTRY
NEWS	15	Jul 30	NETFIRST to be removed from STN
NEWS	16	Aug 08	CANCERLIT reload
NEWS	17	Aug 08	PHARMAMarketLetter(PHARMAML) - new on STN
NEWS	18	Aug 08	NTIS has been reloaded and enhanced
NEWS	19	Aug 19	Aquatic Toxicity Information Retrieval (AQUIRE) now available on STN
NEWS	20	Aug 19	IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS	21	Aug 19	The MEDLINE file segment of TOXCENTER has been reloaded
NEWS	22	Aug 26	Sequence searching in REGISTRY enhanced
NEWS	23	Sep 03	JAPIO has been reloaded and enhanced
NEWS	24	Sep 16	Experimental properties added to the REGISTRY file
NEWS	25	Sep 16	Indexing added to some pre-1967 records in CA/CAPLUS
NEWS	26	Sep 16	CA Section Thesaurus available in CAPLUS and CA
NEWS	27	Oct 01	CASREACT Enriched with Reactions from 1907 to 1985
NEWS	28	Oct 21	EVENTLINE has been reloaded
NEWS	29	Oct 24	BEILSTEIN adds new search fields
NEWS	30	Oct 24	Nutraceuticals International (NUTRACEUT) now available on
STN			
NEWS	31	Oct 25	MEDLINE SDI run of October 8, 2002
NEWS	32	Nov 18	DKILIT has been renamed APOLLIT
NEWS	33	Nov 25	More calculated properties added to REGISTRY
NEWS	34	Dec 02	TIBKAT will be removed from STN
NEWS	35	Dec 04	CSA files on STN
NEWS	36	Dec 17	PCTFULL now covers WP/PCT Applications from 1978 to date
NEWS	37	Dec 17	TOXCENTER enhanced with additional content
NEWS	38	Dec 17	Adis Clinical Trials Insight now available on STN

NEWS 39 Dec 30 ISMEC no longer available

NEWS EXPRESS December 31 CURRENT WINDOWS VERSION IS V6.01a,  
CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),  
AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002  
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NEWS WWW CAS World Wide Web Site (general information)

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\* \* \* \* \* STN Columbus \* \* \* \* \*

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=> file caplus

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FILE 'CAPLUS' ENTERED AT 08:28:03 ON 03 JAN 2003

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FILE COVERS 1907 - 3 Jan 2003 VOL 138 ISS 2

FILE LAST UPDATED: 2 Jan 2003 (20030102/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

=> e wo9608485/pn

E1	1	WO9608483/PN
E2	1	WO9608484/PN
E3	1 -->	WO9608485/PN
E4	1	WO9608486/PN
E5	1	WO9608487/PN
E6	1	WO9608488/PN
E7	1	WO9608489/PN
E8	1	WO9608490/PN
E9	1	WO9608491/PN
E10	1	WO9608492/PN
E11	1	WO9608493/PN
E12	1	WO9608494/PN

=> s e3

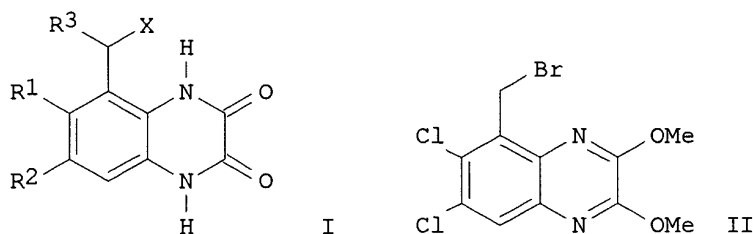
L1 1 WO9608485/PN

=> d ll all

L1 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2003 ACS  
 AN 1996:428452 CAPLUS  
 DN 125:86683  
 TI Preparation of quinoxalinediones as NMDA receptor antagonists  
 IN Mowbray, Charles Eric; Stobie, Alan; Bull, David John; Carr, Christopher  
 Lee; Fray, Michael Johnathan  
 PA Pfizer Limited, UK; Pfizer Research and Development Company, N.V./s.A.;  
 Pfizer Inc.  
 SO PCT Int. Appl., 54 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 IC ICM C07D403-06  
 ICS A61K031-495  
 CC 28-17 (Heterocyclic Compounds (More Than One Hetero Atom))  
 Section cross-reference(s): 1

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9608485	A1	19960321	WO 1995-EP3483	19950901 <--
	W: CA, FI, JP, MX, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	CA 2199845	AA	19960321	CA 1995-2199845	19950901
	EP 781279	A1	19970702	EP 1995-931989	19950901
	EP 781279	B1	20010613		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, NL, PT, SE				
	JP 09511523	T2	19971118	JP 1995-509872	19950901
	JP 2898097	B2	19990531		
	ES 2158126	T3	20010901	ES 1995-931989	19950901
	FI 9701026	A	19970312	FI 1997-1026	19970312
	US 5783572	A	19980721	US 1997-793896	19970312
PRAI	GB 1994-18443	A	19940913		
	WO 1995-EP3483	W	19950901		
OS	CASREACT 125:86683; MARPAT 125:86683				
GI					



AB The title compds. [I; R1, R2 = F, Cl, Br, Me, Et, CF3; R3 = H, Me, Et; X = (substituted) 1,2,4-triazol-1-yl, imidazol-1-yl, pyrazol-1-yl, etc.], useful in the treatment of acute neurodegenerative and chronic neurol. disorders, were prepd. Thus, reaction of quinoxaline II with 1,2,4-triazole in the presence of K2CO3 in AcNMe2 followed by hydrolysis of the intermediate with 2M HCl in dioxane afforded I [R1 = R2 = Cl; R3 = H; X = 1,2,4-triazol-1-yl]. Compds. I are effective at 0.01-1 mg/kg (i.v.).

ST quinoxalinedione NMDA receptor antagonist prepn; nervous system disease degeneration quinoxalinedione prepn; neurotransmitter antagonist quinoxalinedione prepn

IT Neurotransmitter antagonists  
(prepn. of quinoxalinediones as NMDA receptor antagonists)

IT Nervous system  
(disease, degeneration, treatment; prepn. of quinoxalinediones as NMDA receptor antagonists)

IT	178619-22-2P	178619-23-3P	178619-24-4P	178619-25-5P	178619-26-6P
	178619-27-7P	178619-28-8P	178619-29-9P	178619-30-2P	178619-31-3P
	178619-32-4P	178619-33-5P	178619-34-6P	178619-35-7P	178619-36-8P
	178619-37-9P	178619-38-0P	178619-39-1P	178619-40-4P	178619-41-5P
	178619-42-6P	178619-43-7P	178619-44-8P	178619-45-9P	178619-46-0P
	178619-47-1P	178619-48-2P	178619-49-3P	178619-50-6P	178619-51-7P
	178619-52-8P	178619-53-9P	178619-54-0P	178619-55-1P	178619-56-2P
	178619-57-3P	178619-58-4P	178619-59-5P	178619-60-8P	178619-61-9P
	178619-62-0P	178619-63-1P	178619-64-2P	178619-65-3P	178619-66-4P
	178619-67-5P	178619-68-6P	178619-69-7P	178619-70-0P	178619-71-1P
	178619-72-2P				

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
(prepn. of quinoxalinediones as NMDA receptor antagonists)

IT 6384-92-5, NMDA  
RL: BPR (Biological process); BSU (Biological study, unclassified); MSC (Miscellaneous); BIOL (Biological study); PROC (Process)  
(prepn. of quinoxalinediones as NMDA receptor antagonists)

IT 178620-31-0P  
RL: BYP (Byproduct); PREP (Preparation)  
(prepn. of quinoxalinediones as NMDA receptor antagonists)

IT 75-64-9, reactions 89-69-0, 2,4,5-Trichloronitrobenzene 107-59-5, tert-Butyl chloroacetate 109-73-9, n-Butylamine, reactions 110-89-4, Piperidine, reactions 110-91-8, Morpholine, reactions 288-32-4, Imidazole, reactions 288-88-0, 1H-1,2,4-Triazole 594-39-8 4967-77-5,

Methyl 1,2,3-triazole-4-carboxylate 6972-71-0, 4,5-Dimethyl-2-nitroaniline 7170-01-6 7411-16-7 103433-17-6 153504-81-5 153915-05-0 178620-30-9

RL: RCT (Reactant); RACT (Reactant or reagent)

(prepn. of quinoxalinediones as NMDA receptor antagonists)

IT	131885-38-6P	156349-12-1P	178619-73-3P	178619-74-4P	178619-75-5P
	178619-76-6P	178619-77-7P	178619-78-8P	178619-79-9P	178619-80-2P
	178619-81-3P	178619-82-4P	178619-83-5P	178619-84-6P	178619-85-7P
	178619-86-8P	178619-87-9P	178619-88-0P	178619-89-1P	178619-90-4P
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	178619-96-0P	178619-97-1P	178619-98-2P	178619-99-3P	178620-00-3P
	178620-01-4P	178620-02-5P	178620-03-6P	178620-04-7P	178620-05-8P
	178620-06-9P	178620-07-0P	178620-08-1P	178620-09-2P	178620-10-5P
	178620-11-6P	178620-12-7P	178620-13-8P	178620-14-9P	178620-15-0P
	178620-16-1P	178620-17-2P	178620-18-3P	178620-19-4P	178620-20-7P
	178620-21-8P	178620-22-9P	178620-23-0P	178620-24-1P	178620-25-2P
	178620-26-3P	178620-27-4P	178620-28-5P	178620-29-6P	178620-32-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(prepn. of quinoxalinediones as NMDA receptor antagonists)